

Food Safety and Inspection Service Washington, D.C. 20250

JUN 1 2004

Dr. Marc Cornelis Chief Veterinary Officer Institute for Veterinary Inspection Ministry of Social Affairs, Public Health and Environment Blvd du Regent 27 1000 Brussels Belgium

Dear Dr. Cornelis:

The Food Safety and Inspection Service has completed an enforcement audit of Belgium's meat inspection system. The audit was conducted from January 7 through January 16, 2004. Comments from the government of Belgium have been included as an attachment to the final report. Enclosed is a copy of the final audit report.

If you have any questions regarding the audit or need additional information, please contact me by telephone at 202-720-3781, by fax at 202-690-4040, or by email at sally.stratmoen@fsis.usda.gov.

Sincerely,

Sally Stratmoen

Director

International Equivalence Staff
Office of International Affairs

Enclosure

Dr. Marc Cornelis 2

cc:

Belgium-Jan04 Audit

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Luc Devolder, Economic Counselor, Embassy of Belgium
Tony van der haegen, EU Mission to the US, Washington, DC
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FINAL

MAY 1 4 2004

FINAL REPORT OF AN ENFORCEMENT AUDIT CARRIED OUT IN BELGIUM COVERING BELGIUM'S MEAT INSPECTION SYSTEM

JANUARY 7 THROUGH JANUARY 16, 2004

Food Safety and Inspection Service United States Department of Agriculture

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ABBREVIATIONS AND SPECIAL TERMS USED IN THE REPORT

CCA Central Competent Authority (Federal Agency for the Safety of the

Food Chain, or Food Safety Agency)

DG Directorate General

FSA Food Safety Agency

FSIS Food Safety and Inspection Service

VEA European Community/United States Veterinary Equivalence

Agreement

VIC Veterinarian-In-Charge

PCU Provincial Control Unit

PR/HACCP Pathogen Reduction/Hazard Analysis and Critical Control Point

Systems

SSOP Sanitation Standard Operating Procedures

E. coli Escherichia coli

Salmonella Species

1. INTRODUCTION

The audit took place in Belgium from January 7 through January 16, 2004.

An opening meeting was held on January 7, 2004, in Brussels with the Central Competent Authority (CCA). At this meeting, the auditors confirmed the objective and scope of the audit, the auditors' itineraries, and requested additional information needed to complete the audit of Belgium's meat inspection system.

The auditors were accompanied during the entire audit by representatives from the Central Competent Authority (CCA), the Federal Agency for the Safety of the Food Chain, commonly called the Food Safety Agency (FSA), and/or representatives from the provincial and local inspection offices.

2. OBJECTIVE OF THE AUDIT

This was an enforcement audit. The objective was to determine whether Belgium may continue to export meat products to the United States.

In pursuit of the objective, the following sites were visited: The headquarters of the CCA, one provincial office, one laboratory performing analytical testing for *Listeria monocytogenes*, and one meat processing establishment.

Competent Authority Visits			Comments	
Competent Authority	Central	1	Brussels	
	Provincial	1	Hasselt	
Laboratories	1	Liège		
Meat Processing Establishments			Hasselt	

3. PROTOCOL

This on-site audit was conducted in four parts. One part involved visits with CCA officials to discuss oversight programs and practices, including enforcement activities. The second part involved an audit of a selection of records in Belgium's inspection headquarters or provincial offices. The third part involved an on-site visit to the single meat processing establishment that had been eligible to export to the United States during the previous FSIS audit in July 2003 and was delisted during that audit, to determine the adequacy of corrective actions. The fourth part involved a visit to one private laboratory. The National Reference Laboratory for Microbiology was conducting analyses of field samples for the presence of *Listeria monocytogenes*.

Program effectiveness determinations of Belgium's meat inspection system focused on four areas of risk: (1) sanitation controls, including the implementation and operation of Sanitation Standard Operating Procedures (SSOP), (2) animal disease controls, (3) slaughter/processing controls, including the implementation and operation of HACCP

programs, (4) residue controls, and (5) enforcement controls. Belgium's inspection system was assessed by evaluating these five risk areas.

During the on-site establishment visit, the auditors evaluated the nature, extent and degree to which findings impacted on food safety and public health. The auditors also assessed how inspection services are carried out by Belgium and determined if establishment and inspection system controls were in place to ensure the production of meat products that are safe, unadulterated and properly labeled.

In the opening meeting, the auditors explained to the CCA that its inspection system would be audited in accordance with three areas of focus. First, under provisions of the European Community/United States Veterinary Equivalence Agreement (VEA), the FSIS auditors would audit the meat inspection system against European Commission Directive 64/433/EEC of June 1964. This directive has been declared equivalent under the VEA.

Second, in areas not covered by this directive, the auditors would audit against FSIS requirements. FSIS requirements include daily inspection in all certified establishments, the handling and disposal of inedible and condemned materials, species verification, and FSIS's requirements for HACCP and SSOP programs.

Third, the auditors routinely audit against any equivalence determinations that have been made by FSIS. At this time, no equivalence agreements have been made for Belgium under the provisions of the Sanitary/Phytosanitary Agreement.

4. LEGAL BASIS FOR THE AUDIT

The audit was undertaken under the specific provisions of United States laws and regulations, in particular:

- The Federal Meat Inspection Act (21 U.S.C. 601 et seq.).
- The Federal Meat Inspection Regulations (9 CFR Parts 301 to end), which include the Pathogen Reduction/HACCP regulations.

In addition, compliance with the following Community Directive was also assessed:

• Council Directive 64/433/EEC of June 1964 entitled Health Problems Affecting Intra-Community Trade in Fresh Meat

5. SUMMARY OF PREVIOUS AUDITS

Final audit reports are available on FSIS' website at the following address: https://www.fsis.usda.gov/OPPDE/FAR/index.htm.

The following deficiencies were identified during the FSIS audit of Belgium's meat inspection system that was conducted in December 2002:

- Enforcement of U.S. requirements was insufficient: one of the two certified establishments was delisted for failure to meet U.S. requirements.
- SSOP implementation was inadequate in one establishment: there was inadequate documentation of pre-operational and operational sanitation activities, corrective actions, and preventive measures.
- Maintenance of over-product equipment had been neglected (seriously in one establishment, to a minor degree in the other).
- Excessive condensation had not been addressed in one establishment.
- Containers for edible product were not adequately cleaned before use in one establishment.
- There was inadequate separation of containers for edible and inedible product in one establishment.

The following deficiency was identified during the FSIS audit of Belgium's meat inspection system that was conducted in July 2003:

• One establishment was audited and was delisted for failure to provide inspection coverage on Saturdays.

6. MAIN FINDINGS

6.1 Legislation

The auditors were informed that the relevant EC Directive, determined equivalent under the VEA, had been transposed into Belgium's legislation.

The following legal documents provide the legal basis for Belgium's meat inspection service authority:

- Loi du 5 septembre 1952 relative à l'expertise et au commerce des viandes [Law of September 5, 1952, concerning inspection of and trade in meat].
- Arrêté Royal du 30 décembre 1992 relatif à l'agrément et aux conditions d'installation des abattoirs et d'autres établissements [Royal Decree of December 30, 1992, concerning the approval of structural facilities of slaughter and other establishments].
- Arrêté Royal du 4 juillet 1996 relatif aux conditions générales et spéciales d'exploitation des abattoirs et d'autres établissements [Royal Decree of July 4, 1996,

concerning general and specific daily functioning of slaughter and other establishments].

- Arrêté Royal du 9 mars 1953 concernant le commerce des viandes de boucherie et règlement l'expertise des animaux abattus à l'interieur du pays [Royal Decree of March 9, 1953, concerning trade of meat and control of inspection of slaughter animals within the country].
- Arrêté Ministerial du 18 novembre 1991 relatif à l'examen visant à déceler la présence de trichines dans les viandes fraîches provenant l'animaux domestiques de l'espèce porcine, de chevaux et de sangliers ou d'autres espèces de gibier sensibles à la trichinose [Ministerial Decree of November 18, 1991, concerning examination of fresh meat for the presence of trichinosis].

6.2 Government Oversight

The Federal Agency for the Safety of the Food Chain, commonly called the Food Safety Agency (FSA) has four Directorates General: one for Control Policy, one for Control, one for Laboratories, and one for Corporate Services. The Directorate for Control Policy (roughly equivalent to FSIS's Office of Program Development, Policy, and Equivalence) establishes process standards. The Directorate for Control (roughly equivalent to FSIS's Office of Field Operations) carries the responsibility for inspection/audit services and enforcement of process and product standards. This Directorate General (DG) for Control is divided into eleven Provincial Control Units (PCUs), one for each of the 10 Provinces and one for the capital city of Brussels. The DG for Control also has two Coordinators, one for the Flemish-speaking (northern) half of the country and one for the French-speaking (southern) half. These Coordinators supervise the Heads of the PCUs.

6.2.1 CCA Control Systems

When the management of an existing establishment wishes to become eligible to export to the United States, the manager makes an application to the PCU. A Provincial Official Inspector conducts an administrative and technical inquiry and submits a report of the results to the Chief of the PCU, who, in turn, makes a recommendation to the DG Control Headquarters on the basis of the report. The final approval for U.S.-export certification is the responsibility of DG Control. In order to qualify for eligibility to export to the United States, an establishment must first meet EC requirements and must be eligible to produce for inter-community trade. If there is any question regarding the full eligibility of the establishment, a headquarters official from DG Control - Transformation may visit the premises on-site before a final approval is granted.

Communications regarding FSIS requirements are transmitted directly and promptly, by the agricultural section of the American Embassy in Brussels, to the Head of FSA International Affairs [the Counsellor General, DG Control Policy]. He transmits them, as well as other official guidelines and instructions that are issued by DG Control Policy, to the DG for Control (whose office is in the same facility). DG Control forwards them by e-mail and through the mail service promptly to the Head of the PCU. The latter, in turn,

provides them immediately to the Veterinarian-In-Charge (VIC) and her alternate. The entire export manual is also available on FSA's Website. Hard copies of official U.S. requirements, including the U.S. Code of Federal Regulations, Directive 5000.1, and the new RTE Directive, were on hand in both the PCU and the establishment inspection office.

In order to maintain U.S. certification, an establishment must be in 100% compliance with a detailed checklist of FSIS requirements. The officials from the PCU ensure that these requirements continue to be met. If any of the requirements are not met, the PCU correlates with DG Control and U.S. eligibility is revoked by DG Control and the action is reported to International Affairs, DG Control Policy, with immediate notification of FSIS.

6.2.2 Ultimate Control And Supervision

The VIC of the establishment is a full-time FSA employee, who performs the inspection coverage of Est. B-156 on a circuit basis, in addition to coverage of other establishments. There is also a contract-FSA employee, also a qualified veterinarian, who alternates inspection coverage with the VIC and who has had the same inspection training, including numerous recent, documented, official courses in HACCP and SSOP, as the VIC.

There is a clear-cut chain of command from the headquarters of FSA down to the in-plant inspection personnel. The inspection office of the two veterinarians who share the oversight of the establishment is in the city of Hasselt, some six kilometers from the establishment. Their immediate supervisor, the Head of the Provincial [Meat Inspection] Control Unit, or PCU, also has his office in the same facility, so there is almost daily face-to-face correlation between the veterinarians performing the inspection oversight and their immediate supervisor. The direct supervisor of the Head of the PCU (the Chief of the Province) —the Coordinator—has his office in a different building in the same city. The latter is directly supervised by the DG of Control in Brussels. These offices will be co-located in the next several months.

There is a full, written audit/review program with established system controls, including reporting documents and distribution of reports at all levels, as well as documented evidence of daily inspection in the establishment. This documentation was provided.

There are also detailed, written guidelines for supervision of veterinarians and other field FSA employees. Examples were provided. Written reports are required, produced, and distributed to the employees supervised as well as to the supervisors of the employee being evaluated.

Furthermore, there are written criteria for evaluation of the establishment's HACPP programs by the inspection staff. Detailed forms for this evaluation have been developed and are in daily use. Written reports are produced on a regular basis—some daily, some weekly, some monthly—and copies are maintained on record in the inspection files in the establishment. Copies are also routinely reviewed by the supervisors of the in-plant

inspection staff. An on-going summary report of findings has been established, which refers back to specific findings, in order to facilitate the tracking of problems and the occurrence of trends.

6.2.3 Assignment of Competent, Qualified Inspectors

Applicants wishing employment in the FSA must take civil service examinations. Specific additional examinations are prepared and required for veterinarians. The responsibility for the hiring of veterinarians and other inspection employees lies with the Minister of Public Health. The hiring of independent veterinarians (such as the alternate veterinarian providing coverage in Est. B-156) is organized by the Provincial Control Unit. Both federally-recruited and independent veterinarians are required to spend one year in probationary status, during which they are given specific courses in the various aspects of meat inspection, in close coordination with university faculties for veterinary medicine and meat hygiene, and work together with an experienced official inspector. The FSA's Center for training and Education provides continuing training and education for official inspectors.

Both full-time and contract government employees are prohibited by law from performing any private, establishment-paid tasks at an establishment in which they perform official inspection duties. For the full-time government employees, this is regulated in the Law of July 13, 1981, "Creation of an Institute of Veterinary Expertise." A private-practice veterinarian may be hired as a part time or contract government employee, but may not perform any private, establishment-paid tasks in any establishment in which he/she has official duties, nor may he have any additional conflicts of interest. This is regulated by the Royal Decree of July 4, 1986.

There are no other conflict-of-interest concerns with the alternate veterinarian, because (1) no animals slaughtered in Belgium are eligible for use in U.S.-eligible product, (2) her practice does not include swine, and (3) she is even legally forbidden to treat the companion animals of establishment employees. Furthermore, her practice is not in the same community as the establishment.

If either of the veterinarians is unable perform inspection coverage, the other performs the service. They organize vacations in advance so that they are never absent at the same time. There have been no instances in which inspection coverage was not provided due to absence of both of these veterinarians.

There are no budgetary restrictions on the hours of inspection coverage at the establishment. The veterinarians are free to spend as much time on the premises as they feel is necessary.

6.2.4 Authority and Responsibility to Enforce the Laws

Both The VIC and her alternate, as well as all other authorities in the chain of command up to DG Control, have full regulatory authority from retention of product up to and including suspension of operations.

There are thorough written procedures for inspection controls, duties, and activities. Examples were provided. The correct implementation of these programs is ensured by both the Head of the PCU and DG Control Headquarters.

If the establishment management personnel note a microbiological problem involving any product, they are legally required to inform the VIC and to initiate a recall. If the VIC notes a public health concern as a result of export/import inspections, supervisory visits, in-plant inspections, or upper level audits/reviews, she immediately retains the affected product and notifies her supervisor in the PCU for further action. There is also a fully-implemented Rapid-Alert-System in Belgium, as mandated by the European Commission.

6.2.5. Adequate Administrative and Technical Support

FSA has the ability to support a third-party audit. Administrative and Technical Support appeared to be adequate at all levels.

6.3 Headquarters Audit

The auditors conducted a review of inspection system documents. This records review was conducted at the headquarters office of FSA in Brussels and in the provincial office in Hasselt, which provides supervisory oversight for the establishment. The records review focused primarily on food safety hazards and included the following.

- Internal review reports
- Supervisory visits to establishments that were certified to export to the U.S.
- Training records for inspectors and laboratory personnel
- New laws and implementation documents such as regulations, notices, directives and guidelines
- Sampling and laboratory analyses for microbiology
- Sanitation and processing inspection procedures and standards
- Control of inedible and condemned materials
- Export product inspection and control including export certificates
- Enforcement records, including examples of consumer complaints, recalls, seizure and control of noncompliant product, and withholding, suspending, withdrawing inspection services from or delisting an establishment that is certified to export product to the United States.

No concerns arose as a result of the examination of these documents.

7. ESTABLISHMENT AUDITS

The FSIS auditors visited the single meat-processing establishment that had been eligible to export to the United States during the previous FSIS audit in July 2003, and was delisted during that audit, to determine the adequacy of corrective actions taken.

8. RESIDUE AND MICROBIOLOGY LABORATORY AUDITS

During the laboratory audits, emphasis is placed on the application of procedures and standards that are equivalent to United States requirements. Residue laboratory audits focus on sample handling, sampling frequency, timely analysis data reporting, analytical methodologies, tissue matrices, equipment operation and printouts, detection levels, recovery frequency, percent recoveries, intra-laboratory check samples, and quality assurance programs, including standards books and corrective actions.

Microbiology laboratory audits focus on analyst qualifications, sample receipt, timely analysis, analytical methodologies, analytical controls, recording and reporting of results, and check samples. If private laboratories are used to test United States samples, the auditors evaluate compliance with the criteria established for the use of private laboratories under the FSIS Pathogen Reduction/HACCP requirements.

The National Reference Laboratory for Microbiology in the Faculty of Veterinary Medicine at the University of Liège was audited.

No deficiencies were noted.

9. SANITATION CONTROLS

As stated earlier, the FSIS auditors focus on five areas of risk to assess Belgium's meat inspection system. The first of these risk areas that the FSIS auditors reviewed was Sanitation Controls.

Based on the on-site audit of the establishment, Belgium's inspection system had controls in place for SSOP programs, all aspects of facility and equipment sanitation, the prevention of actual or potential instances of product cross-contamination, good personal hygiene and practices, and good product handling and storage practices.

In addition, Belgium's inspection system had controls in place for water records, chlorination procedures, back-siphonage prevention, separation of operations, temperature control, work space, ventilation, welfare facilities, and outside premises.

9.1 SSOP

The establishment was evaluated to determine if the basic FSIS regulatory requirements for SSOP were met, according to the criteria employed in the U.S. domestic inspection program. The SSOP program in the establishment was found to meet the basic FSIS regulatory requirements.

9.2 EC Directive 64/433

In the establishment, the provisions of EC Directive 64/433 were effectively implemented.

10. ANIMAL DISEASE CONTROLS

The second of the five risk areas that the FSIS auditors reviewed was Animal Disease Controls. These controls include ensuring adequate animal identification, control over condemned and restricted product, and procedures for sanitary handling of returned and reconditioned product.

No Belgian slaughter facilities were certified as eligible to export to the U.S. at this time. Therefore, an assessment of Belgium's animal disease controls was not within the scope of this audit.

There had been no outbreaks of animal diseases with public health significance since the last FSIS audit.

11. SLAUGHTER/PROCESSING CONTROLS

The third of the five risk areas that the FSIS auditors reviewed was Slaughter/Processing Controls. The controls include the following areas: ante-mortem inspection procedures, ante-mortem disposition, humane handling and humane slaughter, post-mortem inspection procedures, post-mortem disposition, ingredients identification, control of restricted ingredients, formulations, processing schedules, equipment and records, and processing controls of cured, dried, and cooked products.

The controls also include the implementation of HACCP systems in all establishments and implementation of a generic *Escherichia coli (E. coli)* testing program in slaughter establishments.

11.1 Humane Handling and Slaughter

No Belgian slaughter facilities are certified as eligible to export to the U.S. at this time.

11.2 HACCP Implementation

All establishments approved to export meat products to the United States are required to have developed and adequately implemented HACCP programs. These programs are evaluated according to the criteria employed in the United States domestic inspection program.

The HACCP program was reviewed during the on-site audit of the establishment. The establishment management had adequately implemented the HACCP requirements.

11.3 Testing for Generic E. coli

No Belgian slaughter facilities were certified as eligible to export to the U.S. at this time. Therefore, the establishment was not required to meet the basic FSIS regulatory requirements for generic *E. coli* testing.

11.4 Testing for *Listeria monocytogenes*

The establishment audited was producing ready-to-eat products (pork shoulder picnic hams) that, if the establishment had not been delisted, would be eligible for export to the United States. Since this product is fully cooked in hermetically-sealed plastic pouches and there is no post-lethality exposure to the environment, the requirement to test the finished product for *Listeria monocytogenes* under Directive 10,240.4 does not apply.

11.5 EC Directive 64/433

In the establishment, the provisions of EC Directive 64/433 were effectively implemented.

12. RESIDUE CONTROLS

The fourth of the five risk areas that the FSIS auditors reviewed was Residue Controls. These controls include sample handling and frequency, timely analysis, data reporting, tissue matrices for analysis, equipment operation and printouts, minimum detection levels, recovery frequency, percent recoveries, and corrective actions.

No Belgian slaughter facilities were certified as eligible to export to the U.S. at this time. All meat for products eligible for export to the United States is imported from eligible establishments in the Netherlands.

12.1 EC Directive 96/22

No Belgian slaughter facilities were certified as eligible to export to the U.S. at this time. Therefore, evaluation of the residue testing program was not within the scope of this audit.

12.2 EC Directive 96/23

No Belgian slaughter facilities were certified as eligible to export to the U.S. at this time. Therefore, evaluation of the residue testing program was not within the scope of this audit.

13. ENFORCEMENT CONTROLS

The fifth of the five risk areas that the FSIS auditors reviewed was Enforcement Controls. These controls include the enforcement of inspection requirements and the testing program for *Salmonella*.

13.1 Daily Inspection in Establishments

Inspection was being conducted daily in the processing establishment.

13.2 Testing for Salmonella

No Belgian slaughter facilities were certified as eligible to export to the U.S. at this time. Therefore, evaluation of the *Salmonella* testing program was not within the scope of this audit.

13.3 Species Verification Testing

At the time of this audit, Belgium was required to test product for species verification. Species verification testing was being conducted as required.

13.4 Monthly Reviews

During this audit it was found that monthly supervisory reviews of the establishment were being performed and fully documented.

Monthly internal supervisory reviews of the in-plant inspection oversight are conducted by both the Head of the PCU in which the establishment is situated and another official, a HACCP specialist from the PCU of the city of Brussels, both of whom have had certified training in HACCP, SSOP, and other special export requirements. Each internal review report is delivered to the Chief of the Province, who reviews and signs it, and sends copies to the internal reviewer and the Veterinarian-In-Charge of the establishment. Copies of the monthly review reports for U.S.-eligible establishments are routinely provided to and reviewed by all levels of the chain of command, including the National Implementation Control Unit of DG-Control Headquarters in Brussels. The records are maintained on file for a minimum of three years. Internal reviews are not announced in advance to establishment management. The Veterinarian-In-Charge is informed approximately one day in advance of an internal supervisory review.

13.5 Inspection System Controls

The CCA had controls in place for restricted product and inspection samples, shipment security, including shipment between establishments, and prevention of commingling of product intended for export to the U.S. with product intended for the domestic market.

In addition, controls were in place for the importation of only eligible meat from other counties for further processing.

Lastly, adequate controls were found to be in place for security items, shipment security, and products entering the establishments from outside sources.

14. CLOSING MEETING

A closing meeting was held on January 16, 2003, in Brussels with the CCA. At this meeting, the primary findings and conclusions from the audit were presented by the auditors.

52/8-2/M)

The CCA understood and accepted the findings.

Gary D. Bolstad, DVM International Audit Staff Officer

15. ATTACHMENTS

Microbiology Laboratory Audit Report Individual Foreign Establishment Audit Forms Belgium's Comments on the Draft Report

U.S. DEPARTMENT OF AGRICULTURE FOOD SAFETY AND INSPECTION SERVICE					REVIEW DATE		NAME OF FO	A-1			
INTERNATIONAL PROGRAMS FOREIGN COUNTRY LABORATORY REVIEW					Jan. 8, 2004		National Reference Laboratory for Microbiology				
FOREIGN GOV'T AGENCY Federal Agency For the Safety of the Food Chain (FASFA)					ege, Belg	ium	ADDRESS OF LABORATORY University of Liege, Faculty of Vet. Med. B-4000 Liege, Sart-Tilman				
NAME 0	FREVIEWER Dr. Gary D. Bolstad			FOREIGN OFFII ristel Huy		iport/Ex			rdinator for Micro	Lab	
	Residue Code/Name			Lm							
SAMPLING PROCEDURES	REVIEW ITEMS Sample Handling	ITEM 01	#	A							
	Sampling Frequency	02		A							
	Timely Analyses	03	EVALUATION CODE	A							
	Compositing Procedure	04	ALUATI	0							
	Interpret Comp Data	05	Ē	0							
	Data Reporting	06		A					The state of the s		
ANALYTICAL PROCEDURES	Acceptable Method	07	ODE	A							
	Correct Tissue(s)	08	EVALUATION CODE	A							
	Equipment Operation	09	ALUA	A							
	Instrument Printouts	10	Ē	0							
	Minimum Detection Levels	11		О							
NCE	Recovery Frequency	12		О							
QUALITY ASSURANCE PROCEDURES	Percent Recovery	13	CODE	0							
CEDI	Check Sample Frequency	14	ATIO	A							
ALITA	All analyst w/Check Samples	15	EVALUATION	A							
on'	Corrective Actions	16	Ē	A							
	International Check Samples	17		A							
REVIEW	Corrected Prior Deficiencies	18	EVAL. CODE	A							
OTHER REVIEW		19	AL CODE								
	JRE OF REVIEWER	20	EVAL					DATE	.		
	JRE OF REVIEWER SOLFALY	John	_					Januar	x 8,2004	/	

FOREIGN COUNTRY LABORATORY F	EVIEW	I REVIEW DATE	NAME OF FOREIGN LABORATORY $A\cdot 2$. National Reference Laboratory for Microbiology			
(Comment Sheet)	,	Jan. 8, 2004				
FOREISN GOV'T AGENCY Federal Agency For the Safety of the Food Chain (FASFA)	CITY & COUNTRY Liege, Belgium		ADDRESS OF LABORATORY University of Liege, Faculty of Vet. Med. B-4000 Liege, Sart-Tilman			
NAME OF REVIEWER Dr. Gary D. Bolstad		IAME OF FOREIGN OFFICIAL Drs. Kristel Huysmans, Import/Export, Brigitte Pochet, Coordinator for Micro Lab				
RESIDUE I ITEM NO.	<u> </u>	COMM	ENTS			

No deficiencies were identified.

Foreign Establishment Audit Checklist

1. ESTABLISHMENT NAME AND LOCATION	I 2. AUDIT D	ATE	3. ESTABLISHMENT NO.	4. NAME OF COUNTRY			
N.V. Vleeswarenfabriek Deko Jan. 9, 5. NAME		04	B-156	Belgium			
		E OF AUDITOR(S)		6. TYPE OF AUDIT			
Dr C			olstad	T			
			ary D. Bolstad X ON-SITE AUDIT DOC				
Place an X in the Audit Results block to in		ncompi		nents. Use O if not applicable art D - Continued	e.		
Part A - Sanitation Standard Operating Procedures (SSOP) Basic Requirements		Audit Results		Audit Results			
7. Written SSOP		<u> </u>	33. Scheduled Sample	conomic Sampling	0		
8. Records documenting implementation. 9. Signed and dated SSOP, by on-site or overall authority.			34. Species Testing				
			35. Residue	0			
Sanitation Standard Operating Procedures (SSOP)				- Other Requirements			
Ongoing Requirements				- Other Requirements			
 Implementation of SSOP's, including monitoring of implements. Maintenance and evaluation of the effectiveness of SSOP's 			36. Export				
12. Corrective action when the SSOPs have falled to prevent of			37. Import				
product contamination or adulteration.	inect		38. Establishment Grounds	and Pest Control			
13. Daily records document item 10, 11 and 12 above.			39. Establishment Construction/Maintenance				
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements			40. Light				
14. Developed and implemented a written HACCP plan.			41. Ventilation		1		
15. Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective a	ctions		42. Plumbing and Sewage		!		
16. Records documenting implementation and monitoring of the		~	43. Water Supply				
HACCP plan. 17. The HACCP plan is signed and dated by the responsible establishment individual.			44. Dressing Rooms/Lavato	pries			
			45. Equipment and Utensils	3			
Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements	j		46. Sanitary Operations				
18. Monitoring of HACCP plan.			47. Employee Hygiene		- 		
19. Verification and validation of HACCP plan.			48. Condemned Product Co	entrol			
20. Corrective action written in HACCP plan.	}	}					
21. Reassessed adequacy of the HACCP plan.	;		Part F - Ir	spection Requirements			
22. Records documenting: the written HACCP plan, monitoring of critical control points, dates and times of specific event occurrences.			49. Government Staffing				
Part C - Economic / Wholesomeness			50. Daily Inspection Covera	ge			
23. Labeling - Product Standards			51. Enforcement				
24. Labeling - Net Weights		ŀ			[
25. General Labeling			52. Humane Handling		0		
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Mo	isture)		53. Animal Identification		0		
Part D - Sampling Generic <i>E. coli</i> Testing			54. Ante Mortem inspection		, 0		
27. Written Procedures		0	55. Post Mortem Inspection		1		
28. Sample Collection/Analysis	·	0			0		
29. Records		0	Part G - Other Regul	latory Oversight Requirements			
Salmonella Performance Standards - Basic Requir	rements	5	66. European Community Din	ectives			
30. Corrective Actions		0	57. Monthly Review		1		
31. Reassessment		0 5	58.		i i		
32. Written Assurance	1	0 5	59.				

60. Observation of the Establishment

B-2

Jan. 9, 2004, Est. B-156, N.V. Vleeswarenfabriek Deko, Hasselt, Belgium.

No deficiencies were found.

NOTE: All previously identified deficiencies had been adequately addressed and corrected.



DG CONTROL POLICY

INTERNATIONAL AFFAIRS

Flie handled by:

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Your references

Our references

Annexes

Office of International Affairs

International Equivalence Staff

date

February 25, 2004

Your letter

PCCB/\$4/\$H\$/\$79 77

USDA - FSIS Sally Stratmoen

Director

30 04 2004

Concern: remarks / draft final audit report

Dear Ms. Stratmoen,

The Food Safety and Inspection Service (FSIS) has completed an enforcement audit of Belgium's meat inspection system. The audit was conducted from January 7 through January 16, 2004. A copy of the draft final audit report was sent to our services.

Hereby we sent you the comments regarding the information in the report:

- 1) Page 5, point 2, 4th line and the boxes: Objective of the audit:
 - to be changed (4th line): CCA, one provincial office, one laboratory performing
 - to be deleted: 4th box (local, 1, Hasselt)
- 2) Page 5, point 3: Protocol; page 6, point 4: Legal basis for the audit; page 13, point 9.2 EC Directive 64/433 and page 14, point 11.5 EC Directive 64/433:

These points refer to Directive EC 64/433 (fresh meat) whereas the establishment Deko NV falls under the Directive EC 77/99 (meat products). This remark is also made on the entrance conference and on the closing meeting as well as on the "Final report of an audit carried out in Belgium covering Belgium's meat inspection system (July 15 through July 28, 2003)".



3) Page 9, point 6.2.2:Ultimate control and supervision:
"Head of the meat inspection headquarters of FSA" is an old term and should be better replaced by "the headquarters of FSA".

Thank you for including these comments as an attachment to the final report.

Sincerely,

Ir. G. Houins

Director-generaal

Cc: Dr. J.-M. DOCHY, director-general, DG Control Embassy of Belgium, Washington